

Figure 1

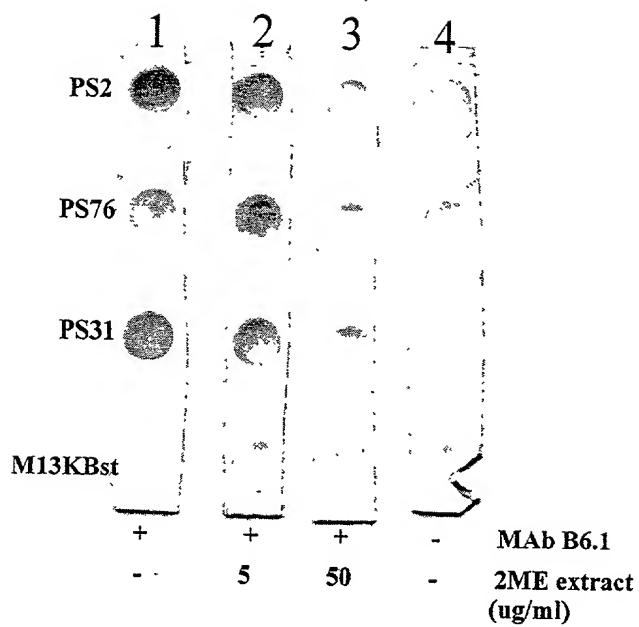


Figure 2

### Reactivity of MAb B6.1 against PS76-carrier protein conjugates

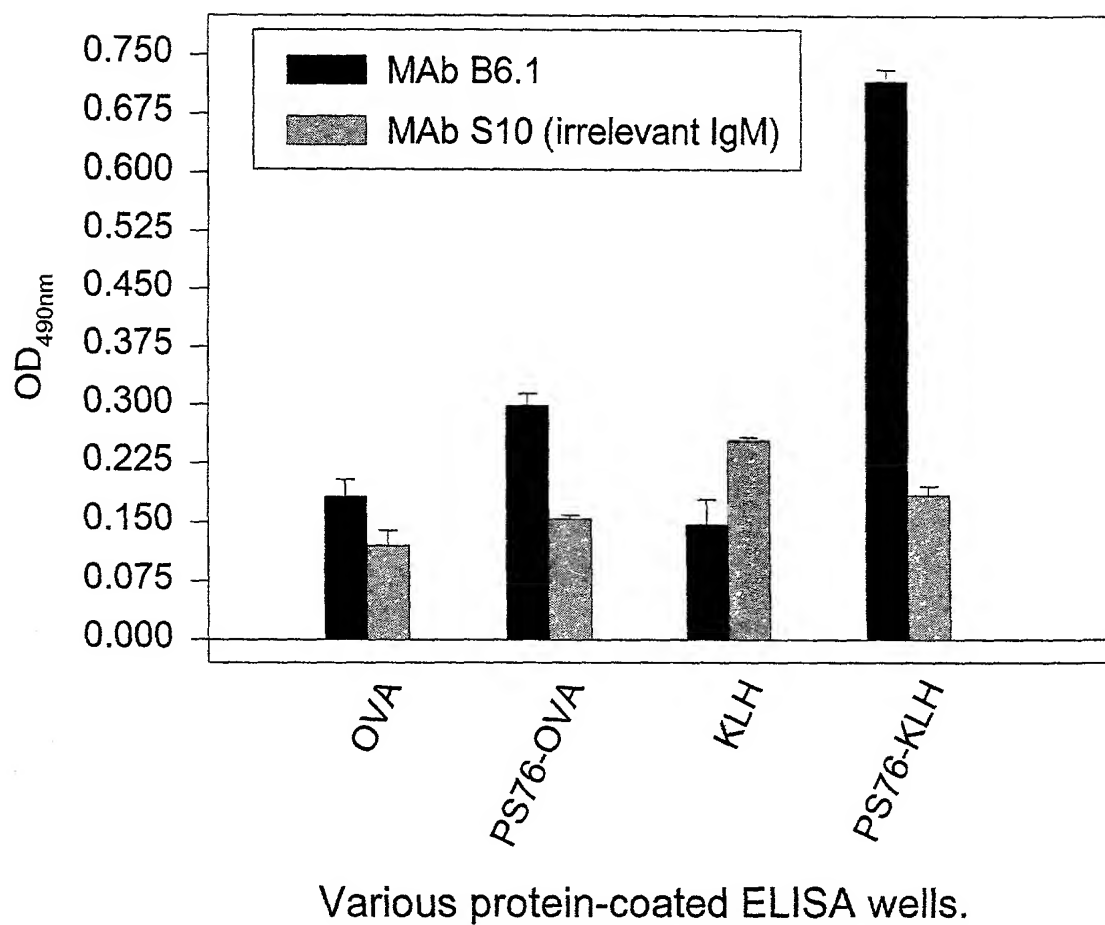


Figure 3

***C. albicans* mannan inhibits MAb B6.1 binding to PS76-KLH conjugate**

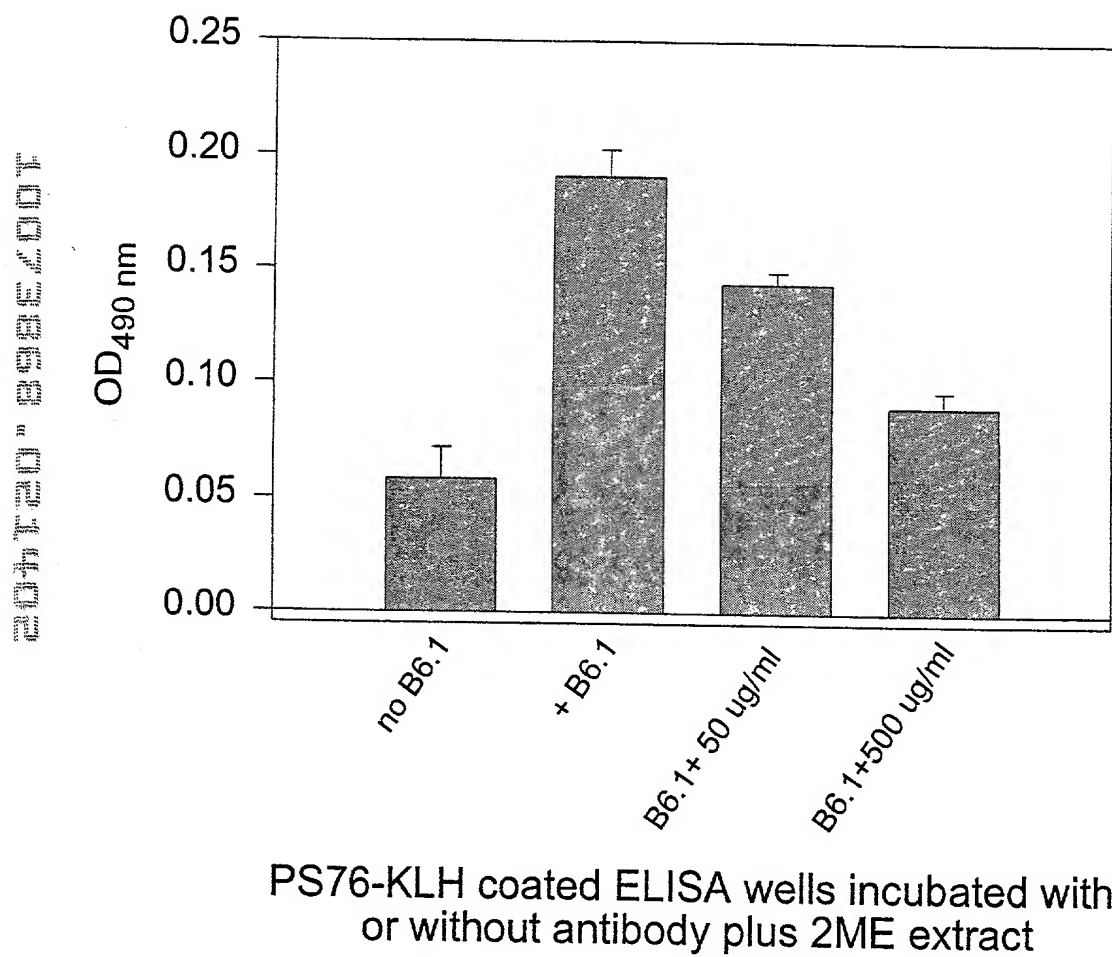


Figure 4

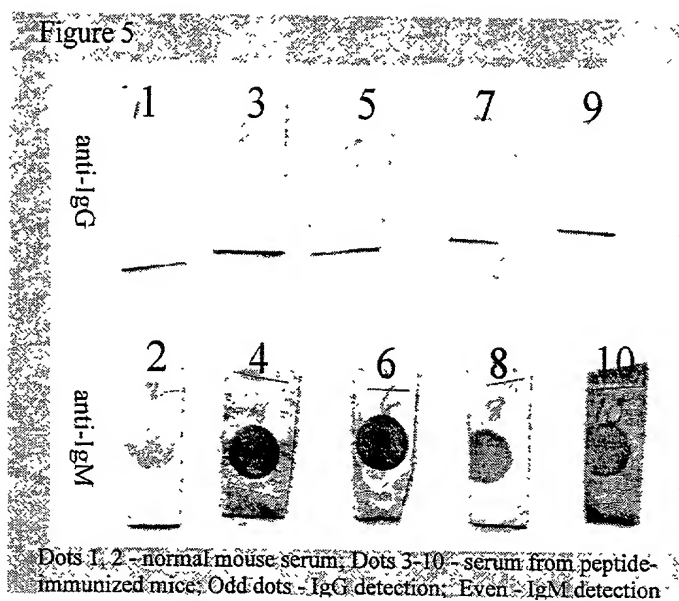
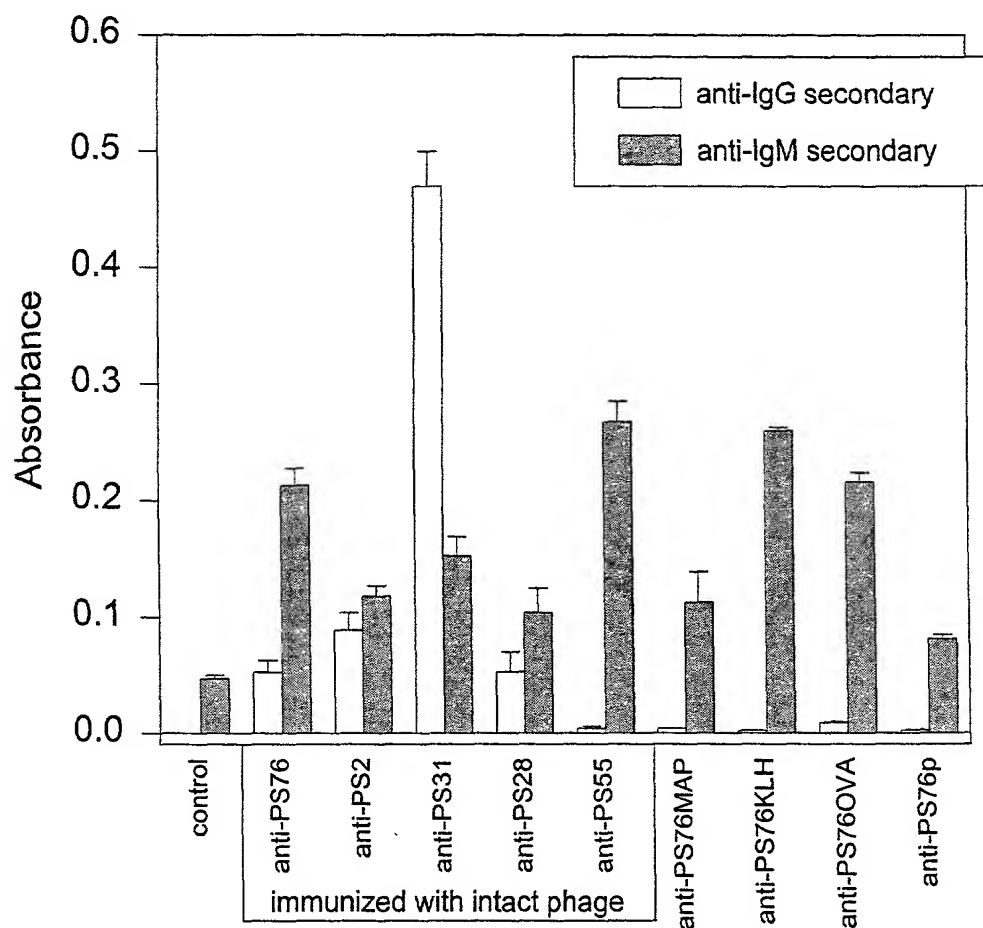


Figure 6

# **Immunization with peptide mimotopes induces anti-*Candida* carbohydrate antibodies in mice**



$$\text{Absorbance} = \text{OD}_{490\text{nm}} (\text{Ab} + 2\text{ME extract}) - \text{OD}_{490\text{nm}} (\text{Ab} + \text{irrelevant CHO})$$

Microtiter plate wells were coated with 2ME extract or an irrelevant carbohydrate. Mice were immunized with synthetic peptide, branched synthetic peptide, peptide carrier protein conjugates, or phage-displayed peptide mimotopes.

1007363040

### Gene 3 protein of phage clone PS76

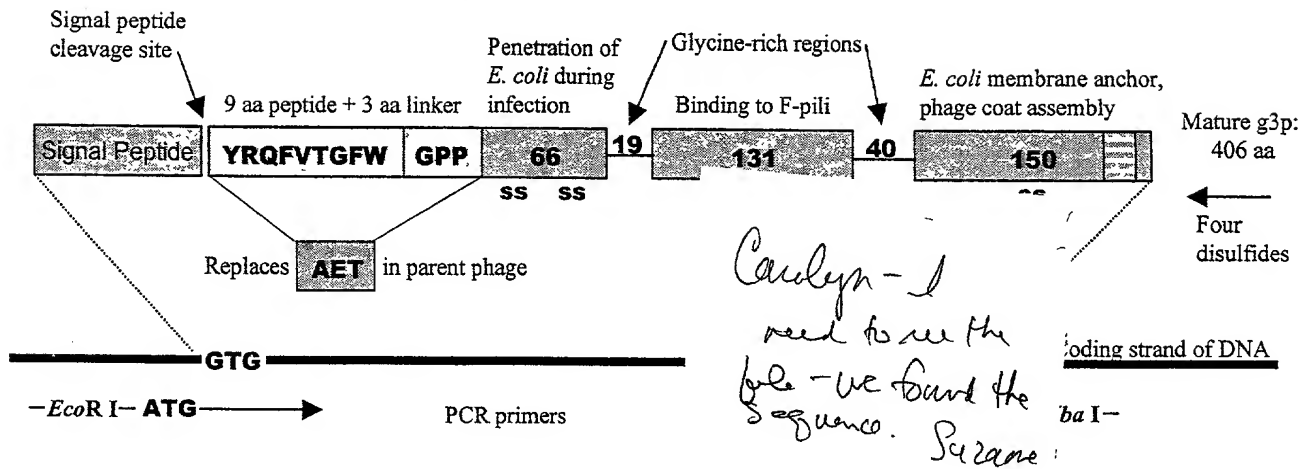


Figure 7

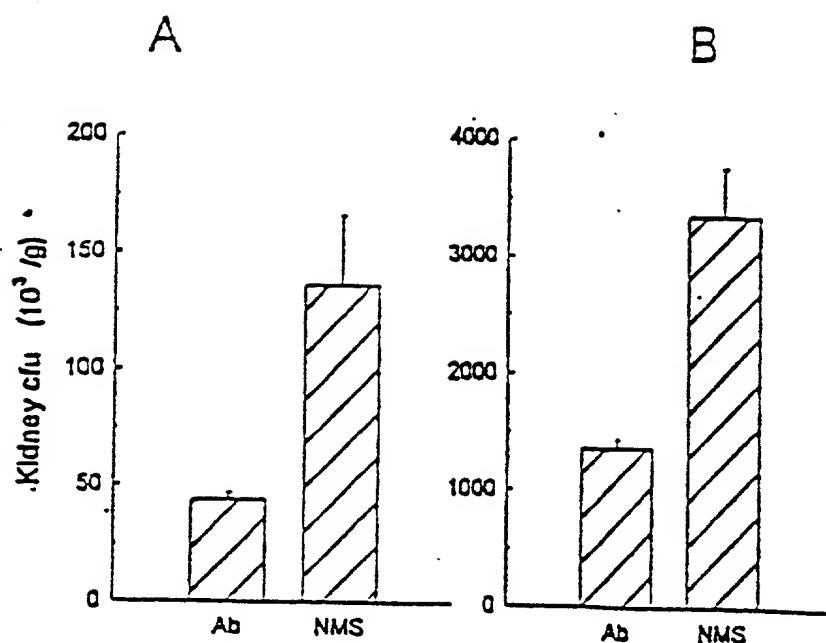
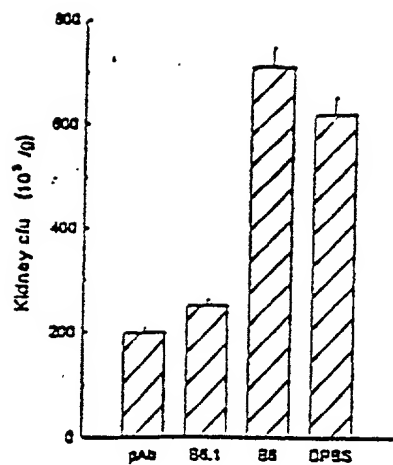


Fig 8.





B

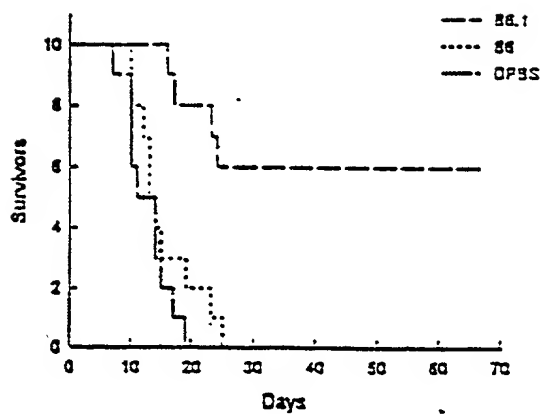


Fig 29

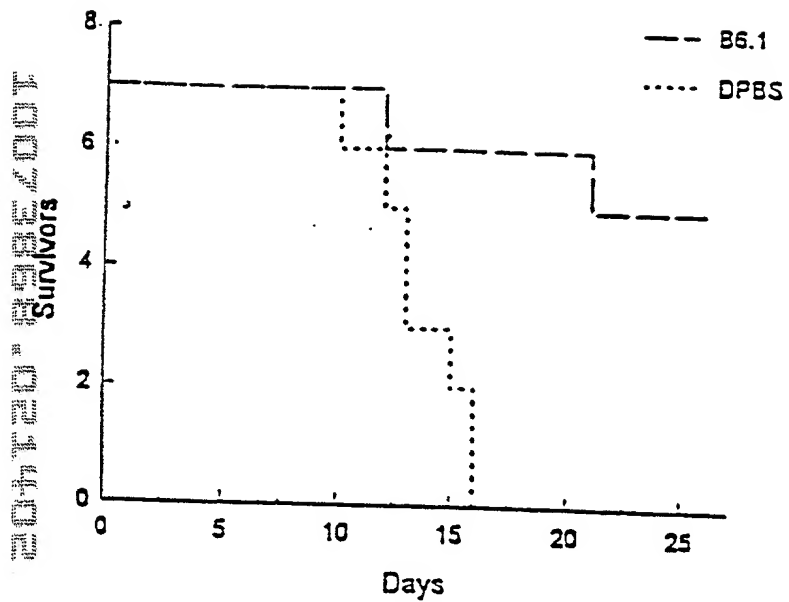


Fig 10

Disseminated Candidiasis By Survival Time  
measurements. Therapeutic effect of MAb B6.1 on candida  
infected mice (one hour infection)

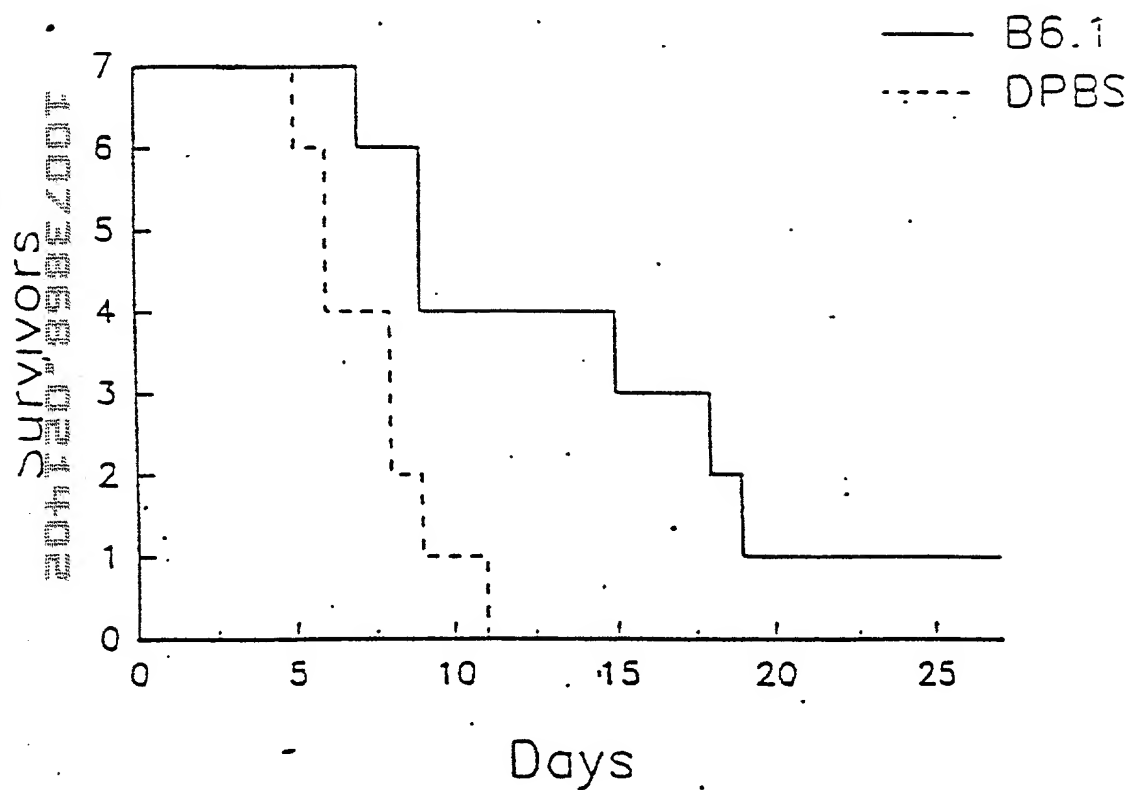


Fig 4  
11

Therapeutic effect of MAb B6.1 on candidal infected mice  
(one hour infection)

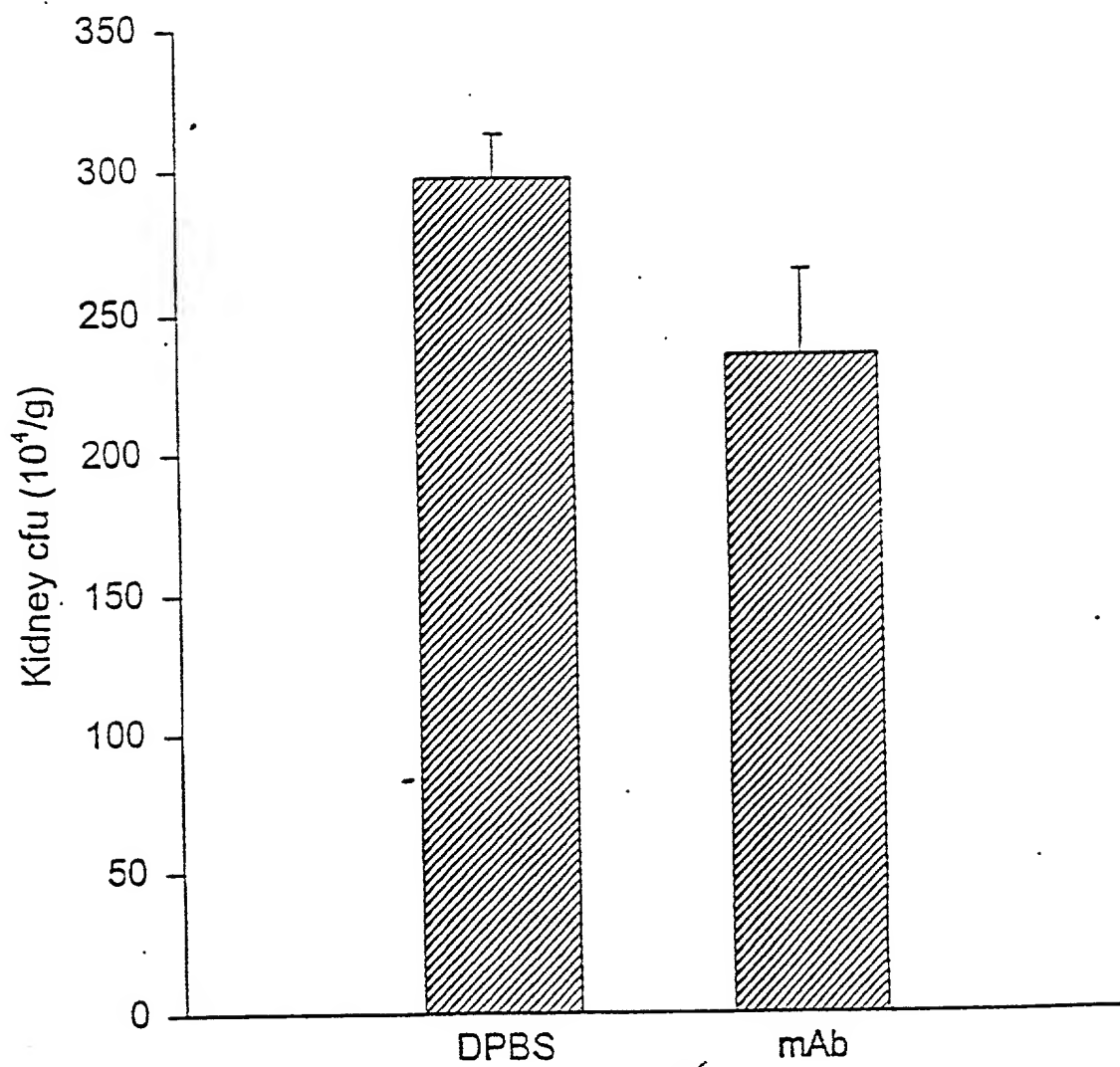


Fig 8  
12

Prophylactic effect of MAb B6.1 on mice to vulvovaginal candidiasis

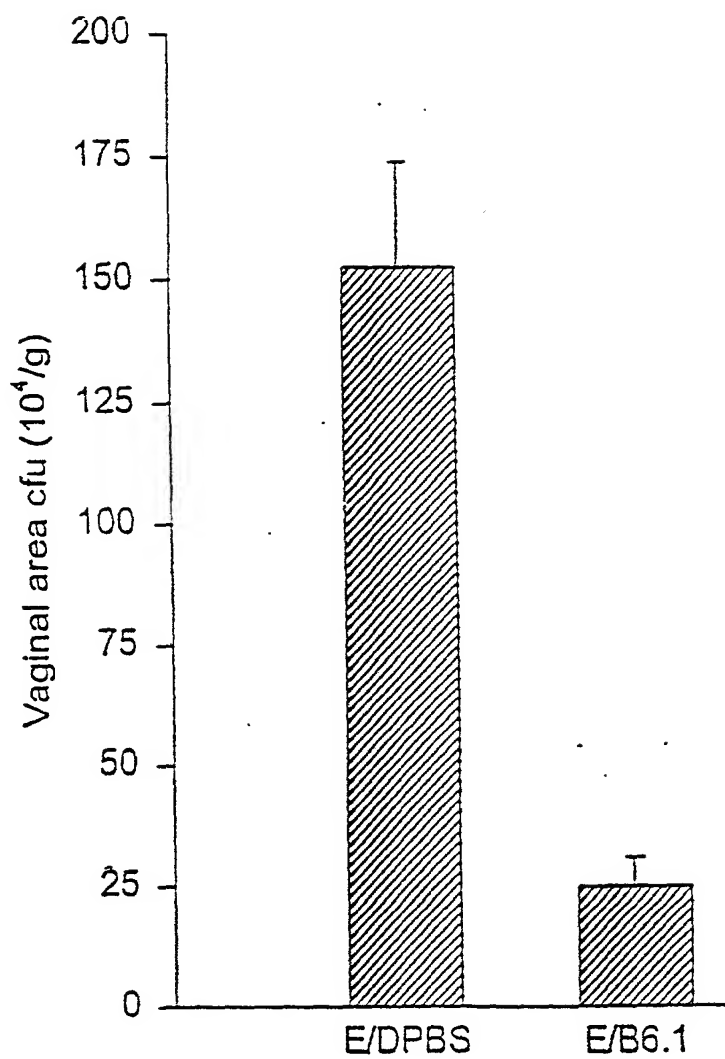


Fig 6  
13

Prophylactic effect of candidal MAbs on mice to  
vulvovaginal candidiasis

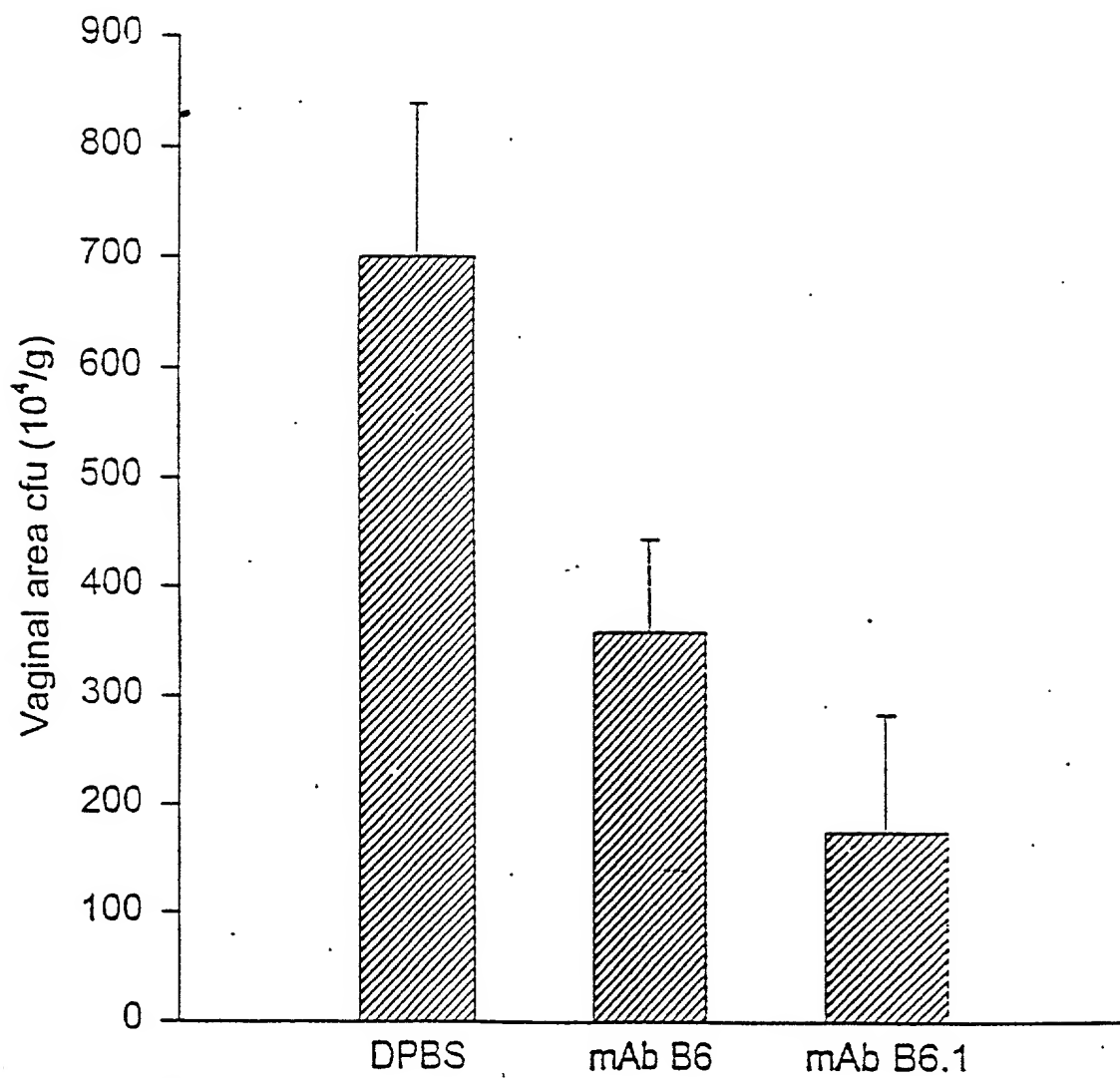


Fig 7.  
14

Effect of active immunization with L-adhesion to mice  
against vulvovaginal candidiasis

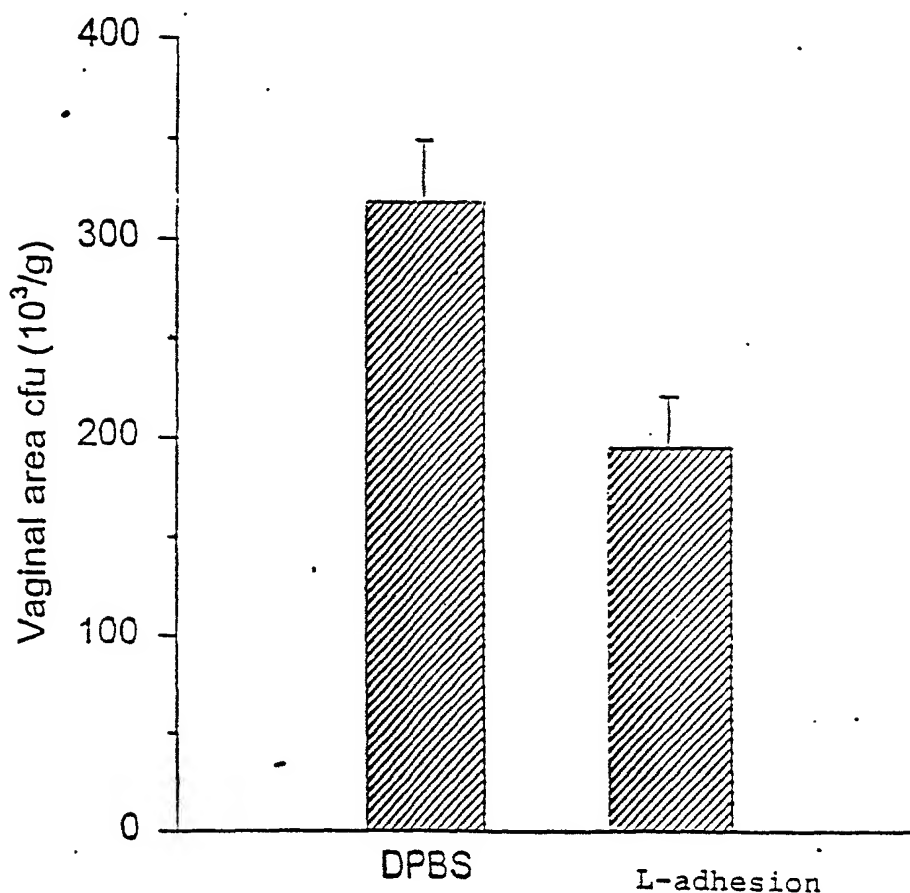


Fig 8/15

L = liposome

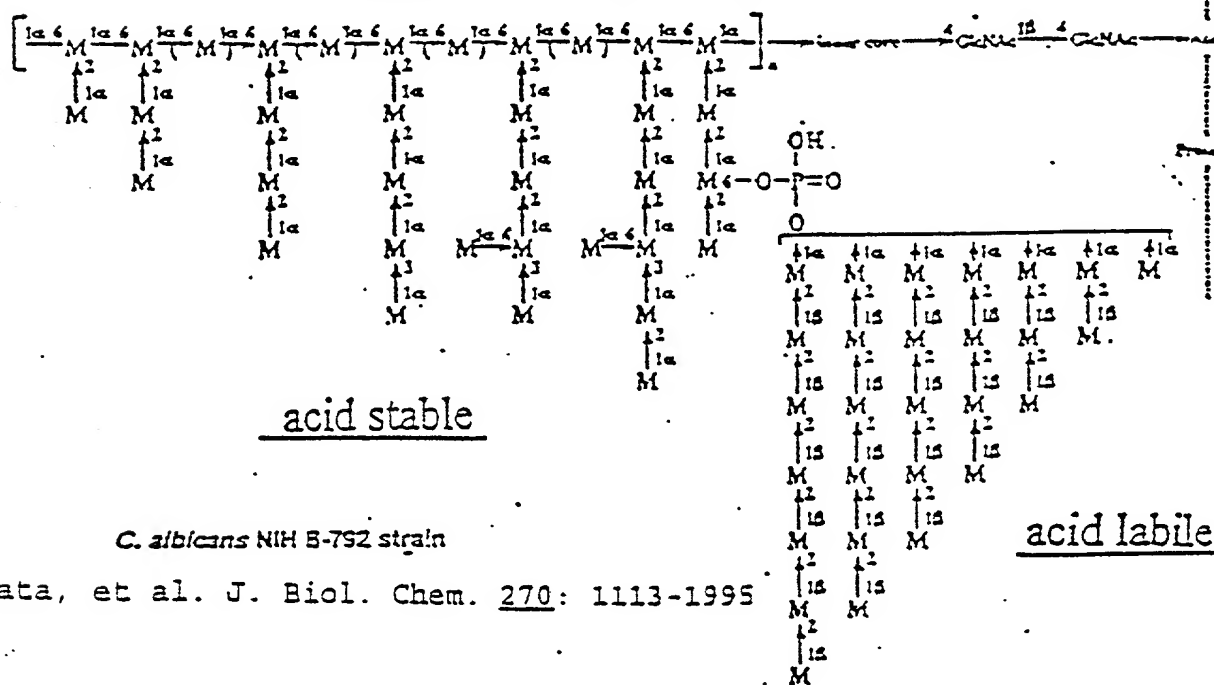
L02ME = liposome-2ME vaccine prep. Animals received 0.2 ml i.v. (178 <sup>µg</sup> 2ME in 0.2ml) weekly for 5 weeks. Estiadiol was given subcu, 72 h later C. albicia ( $5 \times 10^5$ ) gives intravaginally, 48 h after infection vaginal cfu determined.

20073868.024400

Proposed structure of the phosphonennan complex (PMC)-in this case, n-linkage to cell wall protein is shown.

Fig 9.16

### The Phosphomannoprotein Complex



*C. albicans* NIH B-792 strain

Shibata, et al. J. Biol. Chem. 270: 1113-1995



Absorbance at 490 nm

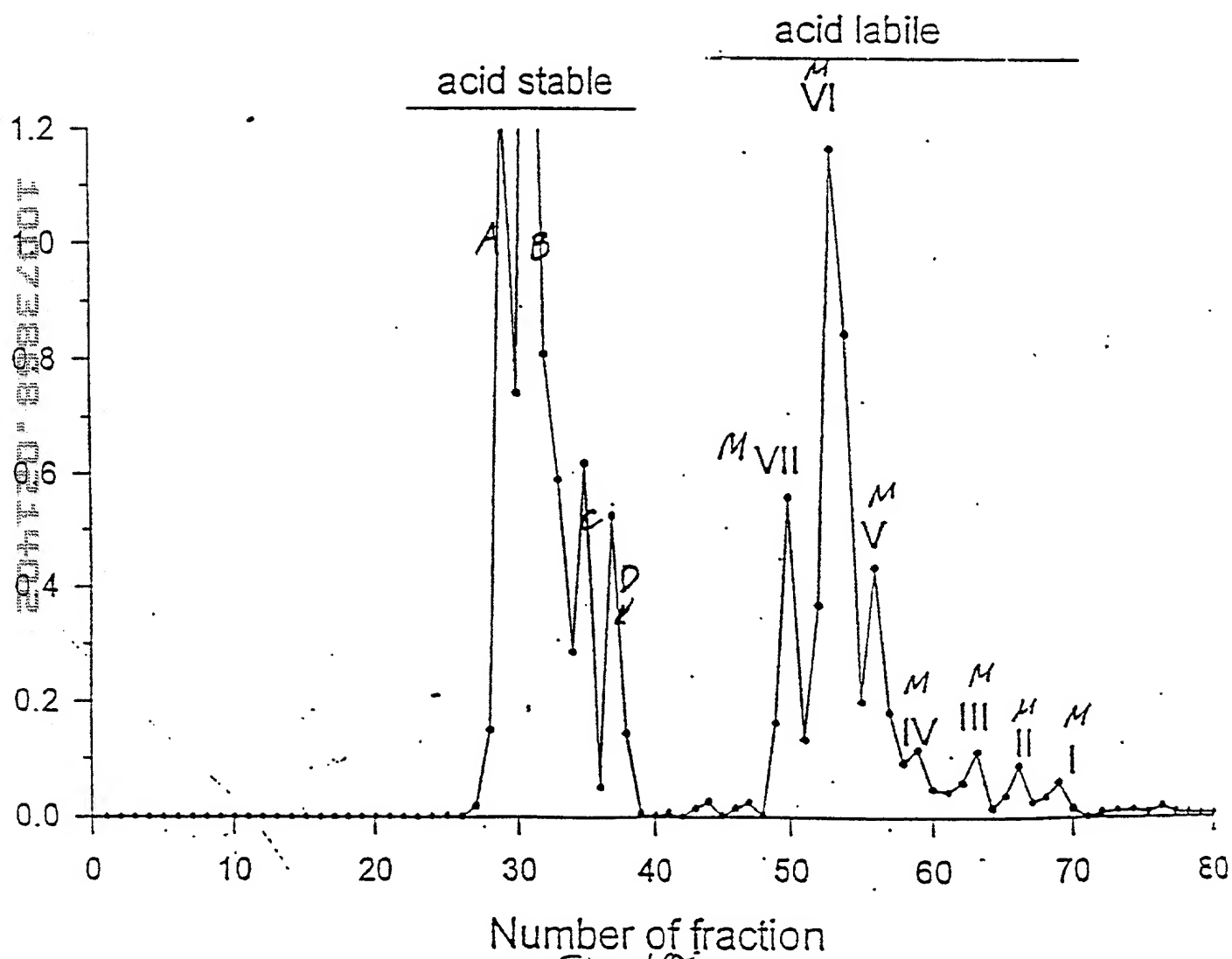
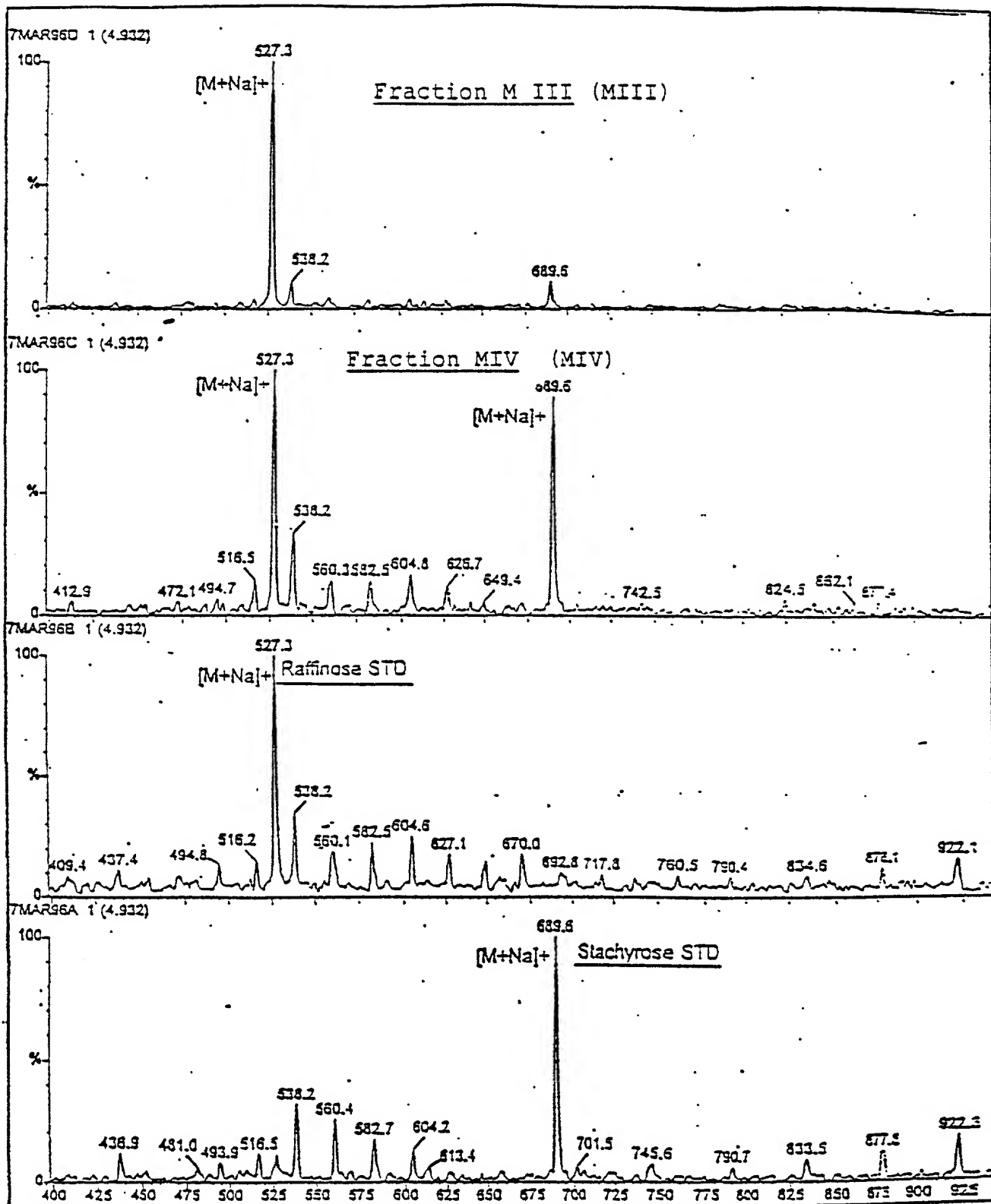


Fig. 10  
17

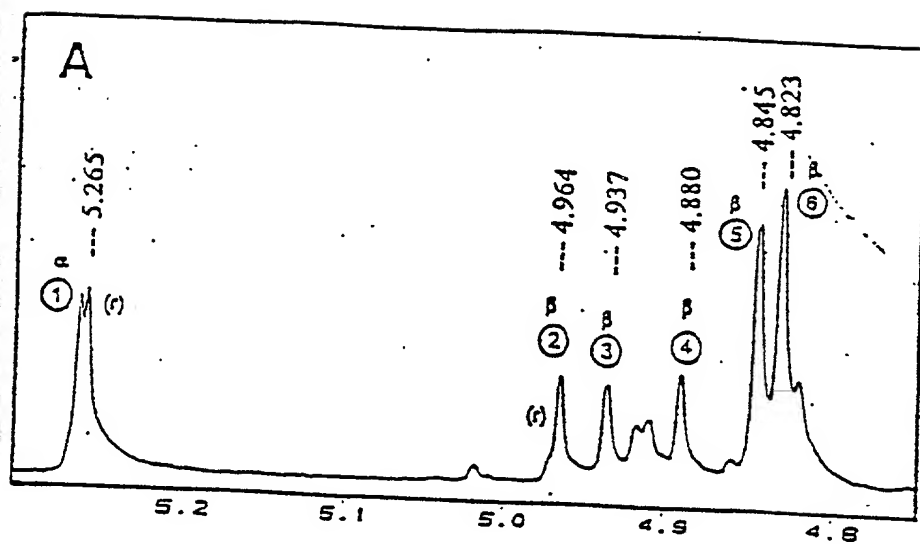
2017-20-09-001



Mass spec results

Fig 11  
18

204720" 238E/001



ppm

Fig 13  
19

dimension H-nmv of B6.1 epitope

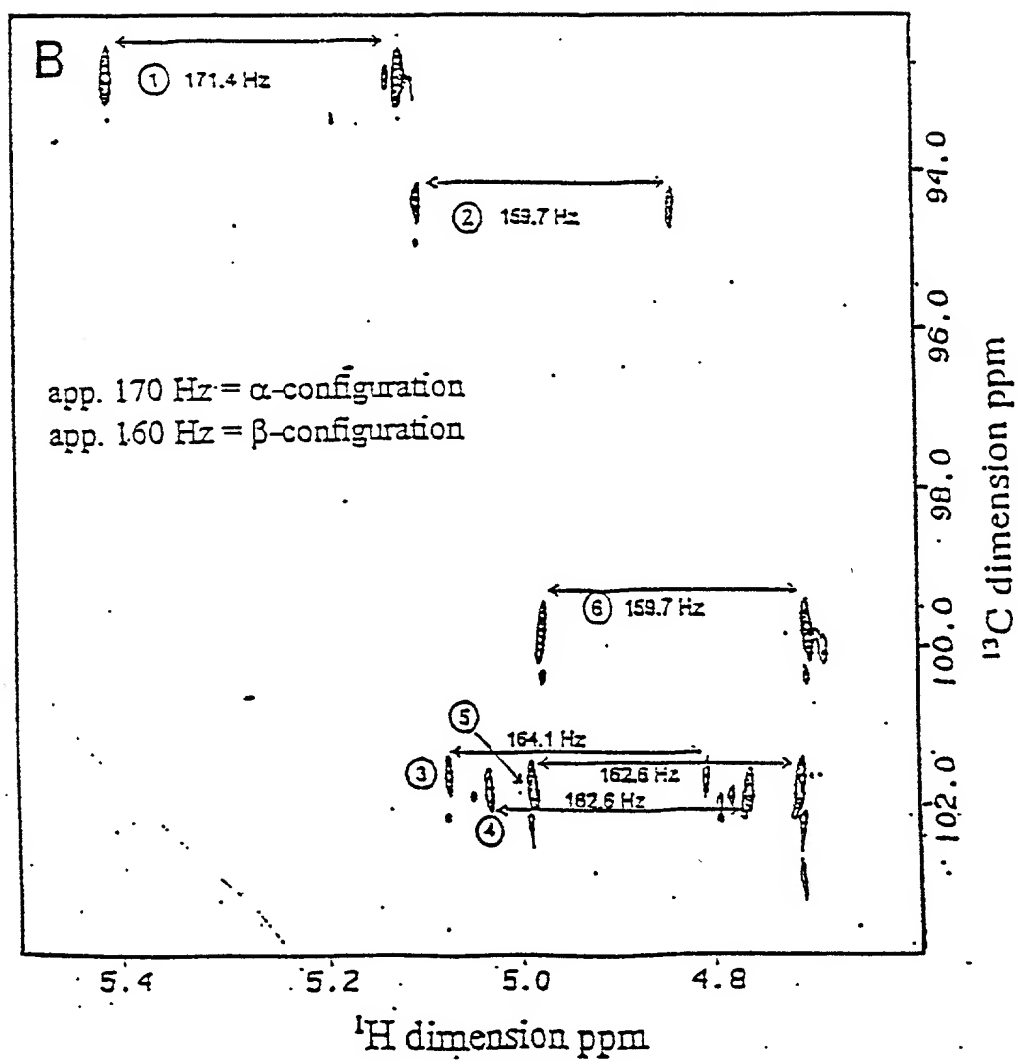


Fig 13

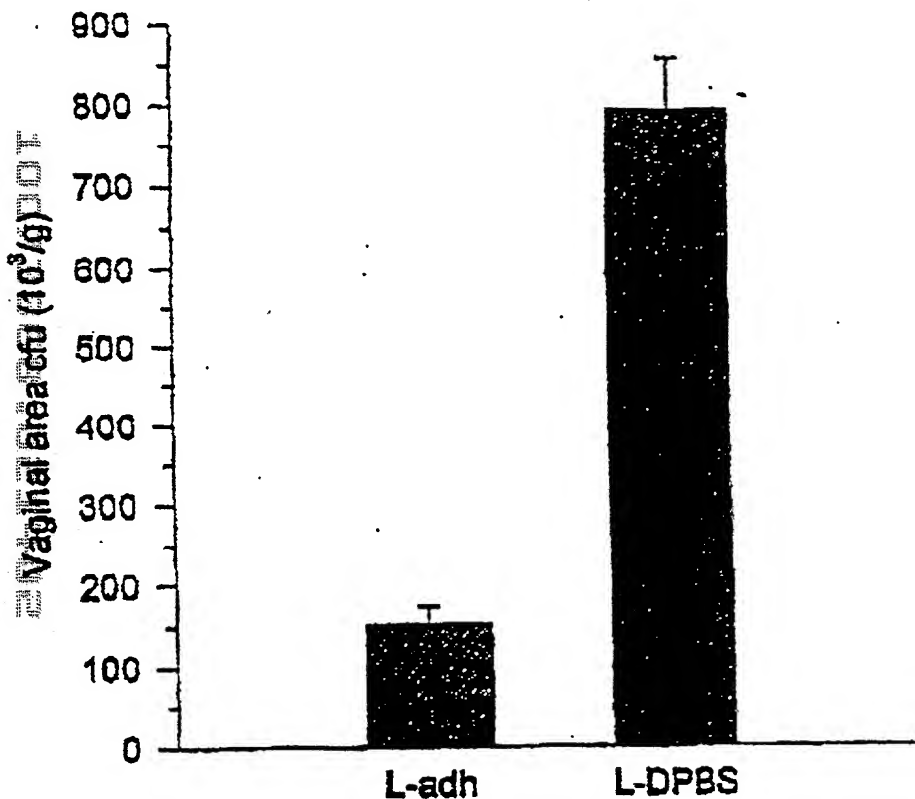
2-DNMR of B6.1 epitope

20

Figure 21

file name: 110596

Therapeutic effect of liposome-2ME (L-adh) as a vaccine source on mice\* against vulvovaginal candidiasis.



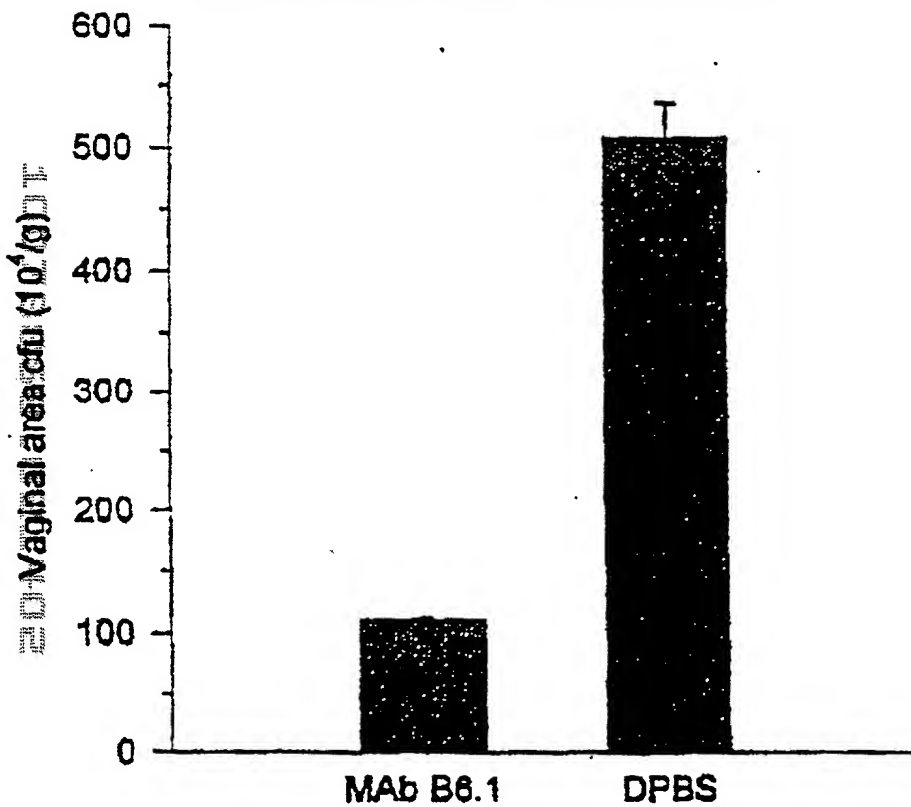
\*Animals were infected with *C. albicans* one hr before  
~~antibody treatment~~ Vaccination

\*\* Mice were intravaginally infected with *Candida albicans* ( $5 \times 10^5$ /mouse) 1 hr before i.v. vaccine treatment. Seven days after the infection, vagina areas were collected, and cfu in the areas were measured.

Fig 15 22

file name: 091396A

Therapeutic effect of MAb B6.1 on mice\*  
against vulvovaginal candidiasis.



\*Animals were infected with *C. albicans* one hr before antibody treatment.

T=0  $5 \times 10^5$  / 10  $\mu$ l IVq.

T=1h MAb B6.1 30  $\mu$ l undiluted 1280, 3.5 mg/ml

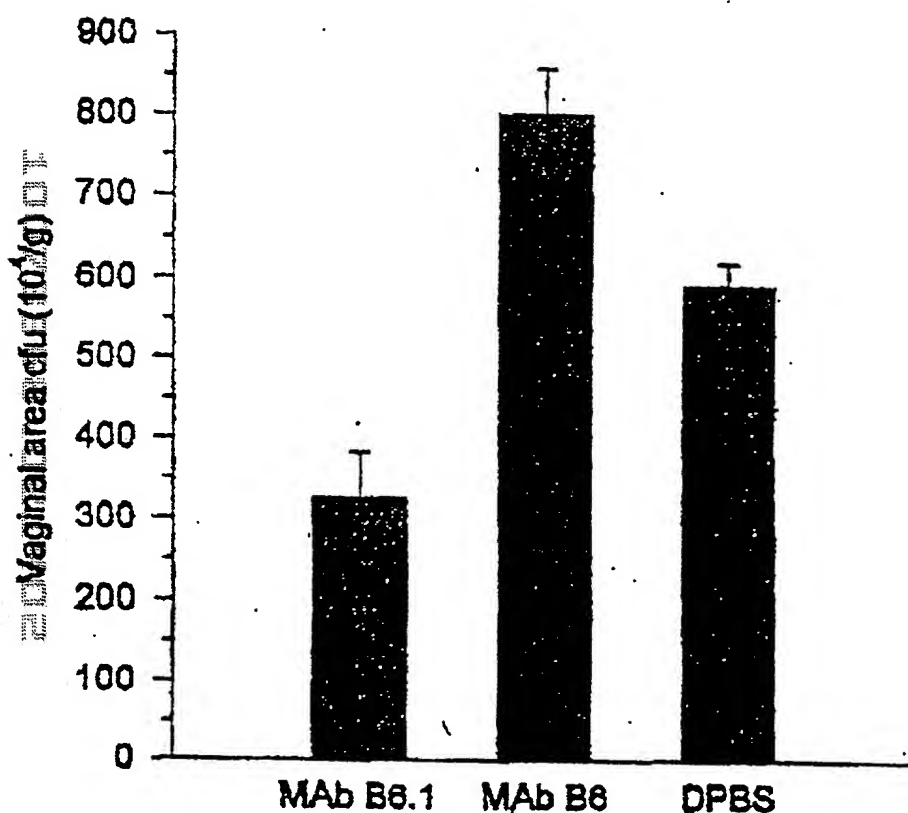
T=24h " " 10  $\mu$ l

T=48h cfc

Fig. 23

file name: 101296

Therapeutic effect of MAb B6.1 & B6 on mice\*  
against vulvovaginal candidiasis.



\*Animals were infected with *C. albicans* four hrs before first antibody treatment. 7/4

T=0: infect  $5 \times 10^5$  i.v.g

T=4h MAb 30  $\mu$ l i.v.g

T=24h " 10  $\mu$ l

T=48h cfu

MAb B6.1  
1280, 3.5mg/ml

MAb B6  
640, 3.5mg/ml

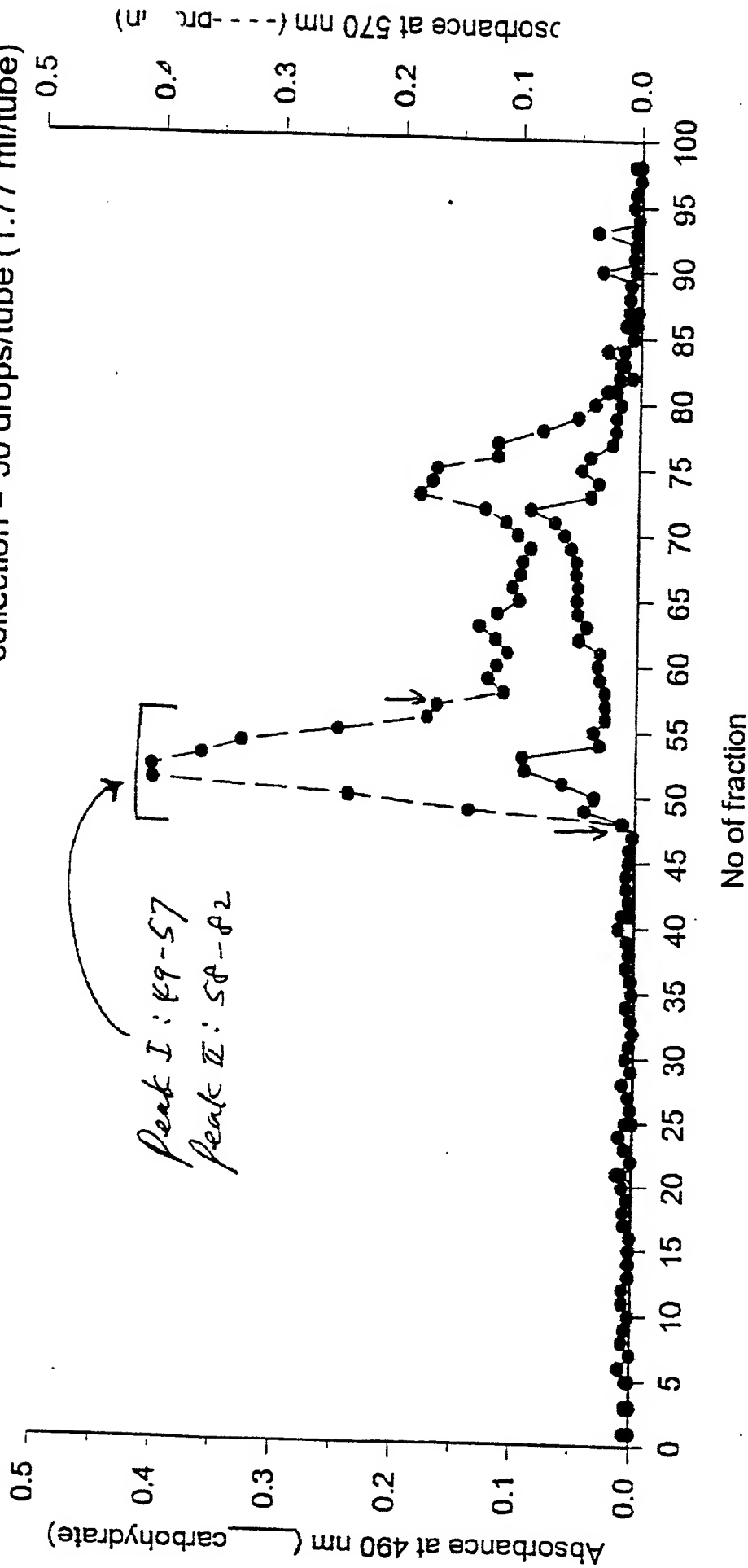
File name: 100296

2011.02.20 09:58:00

Purification of 2ME-BSA by a sephacryl-S-300 column

(Fig. 17)<sup>2</sup>

void volume = 80 ml  
flow rate = 16 ml/hr  
collection = 50 drops/tube (1.77 ml/tube)





File name: 101196a

Purification of 2ME-BSA by a sephacryl S-300 column

(Fig. 78) 29

void volume = 80 ml

flow rate = 16 ml/hr

collection = 50 drops/tube (1.77 ml/tube)

